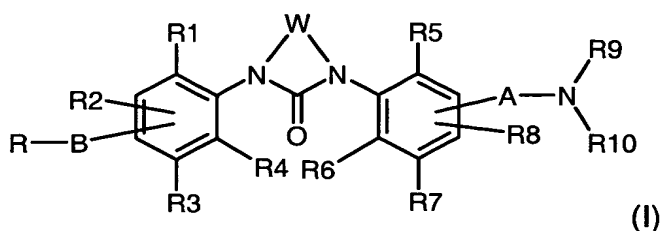


What is claimed is:

1. A compound of formula (I)



wherein

R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or spirocyclic ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R₁₁)(R₁₂), N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R₁₅)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ independently are H or (C₁-C₆)-alkyl;

B is a bond or a linker comprising one or two radicals selected from the group consisting of (C(R₁₉)(R₂₀))_i, C(OR₂₁)(R₂₂), O, N(R₂₃), S, SO, SO₂, and CO;

i is 1, 2 or 3;

R₁₉, R₂₀, R₂₁, R₂₂ and R₂₃ independently are H, (C₁-C₆)-alkyl or aryl;

R₁, R₂, R₃, and R₄ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-

C₈)-alkylene-aryl, S-aryl, N(R₂₄)(R₂₅), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₂₆)(R₂₇), N(R₂₈)CO(R₂₉), N(R₃₀) SO₂(R₃₁) or CO(R₃₂);

R₂₄, R₂₅, R₂₆, R₂₇, R₂₈ and R₃₀ independently are H or (C₁-C₆)-alkyl;

R₂₉, R₃₁, and R₃₂ independently are H, (C₁-C₆)-alkyl or aryl;

W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

n is 2, 3, 4 or 5;

R₅, R₆, R₇ and R₈ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R₃₃)(R₃₄), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₃₅)(R₃₆), N(R₃₇)CO(R₃₈), N(R₃₉) SO₂(R₄₀), CO(R₄₁) or a 5- to 7-membered heterocycle having 1 to 4 heteroatoms selected from the group consisting of O, N and S;

R₃₃ and R₃₄ independently are H or (C₁-C₆)-alkyl, or

R₃₃ and R₃₄ form together with the nitrogen atom to which they are bonded a 5- or 6-membered ring wherein when R₃₃ and R₃₄ form together with the nitrogen to which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring optionally is O or S;

R₃₅, R₃₆, R₃₇ and R₃₉ independently are H or (C₁-C₆)-alkyl;

R₃₈, R₄₀ and R₄₁ independently are H, (C₁-C₆)-alkyl or aryl;

A is a chain $-(C(R42)(R43))_m-$ wherein 0 to 2 members of the chain are optionally replaced by an element selected from the group consisting of O, S, N(R44), CO and SO₂;

5 m is 0, 1, 2, 3, 4 or 5;

R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

10 R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o$ -R45, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl, CO-(CH₂)_o-R45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl, CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

15 R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 4 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

20

R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

o is 0, 1, 2, 3, 4, 5 or 6;

25

R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or more heteroatoms selected from the group consisting of N, O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R51)(R52), SO₂-CH₃ or COOH; or

30

a pharmaceutically acceptable salt of the compound of formula I.

5 2. The compound according to claim 1 wherein

10 R is (C₁-C₆)-alkyl, (C₀-C₂)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₂-C₆)-alkenyl, (C₅-C₈)-cycloalkenyl, (C₇-C₈)-bicycloalkenyl, (C₂-C₆)-alkynyl or a 3- to 7-membered ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 7-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, CON(R₁₁)(R₁₂), N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl or N(R₁₅)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

15 B is a bond O, S, SO₂, CO, OCH(R₂₀), N(R₂₃), CH₂ or CH₂CH₂;

20 R₂₀ and R₂₃ independently are H or (C₁-C₆)-alkyl;

25 R₁, R₂, R₃, and R₄ independently are H, F, Cl, Br, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₀-C₂)-alkylene-aryl, O-(C₀-C₂)-alkylene-aryl, N(R₂₄)(R₂₅), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₂₆)(R₂₇), N(R₂₈)CO(R₂₉) or CO(R₃₂);

30 R₂₉ and R₃₂ independently are H, (C₁-C₆)-alkyl or aryl;

35 n is 2, 3 or 4;

40 R₅, R₆, R₇ and R₈ independently are H, F, Cl, Br, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₀-C₂)-alkylene-aryl, O-(C₀-C₂)-alkylene-aryl, COO-(C₁-C₆)-alkyl or CO(R₄₁);

45 R₄₁ is (C₁-C₆)-alkyl or aryl;

A is a chain $-(C(R42)(R43))_m-$ wherein 1 to 2 members of the chain are optionally replaced by an element selected from the group consisting of O, N(R44) and CO;

5 m is 3 or 4;

R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o-R45$, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl or CO-(CH₂)_oR45; or

10 R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 2 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-C₂)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

 o is 0, 1, 2, 3 or 4;

20 R45 is OH, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or two heteroatoms selected from the group consisting of N, O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₀-C₂)-alkylene-aryl, O-(C₀-C₂)-alkylene-aryl, N(R51)(R52), SO₂-CH₃ or COOH.

25

3. The compound according to claim 2 wherein

 R is (C₁-C₆)-alkyl, (C₃-C₈)-cycloalkyl, or a 5- to 6-membered mono- or bicyclic
30 ring optionally containing one or two heteroatoms selected from the group consisting of N, O, and S, wherein the 5- to 6-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl or O-(C₁-C₆)-alkyl;

B is a bond, O, S, CO OCH₂, N(R₂₃) or CH₂;

R₂₃ is H or (C₁-C₆)-alkyl;

5 R₁, R₂, R₃, and R₄ independently are H, F, Cl, Br, CF₃, O-(C₁-C₆)-alkyl or (C₁-C₆)-alkyl;

W is -CH=CH- or -N=CH-;

10 R₅, R₆, R₇ and R₈ independently are H, F, Cl, Br, OH, CF₃, NO₂, CN, OCF₃ or O-(C₁-C₆)-alkyl;

A is a chain -(C(R₄₂)(R₄₃))_m- wherein 1 member of the chain is optionally replaced by an element selected from the group consisting of O and N(R₄₄);

15 R₉ and R₁₀ independently are H, (C₁-C₈)-alkyl, -(CH₂)_o-R₄₅ or CO-(C₁-C₈)-alkyl; or

R₉ and R₁₀ form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 2 additional heteroatoms selected from the group consisting of O, N and S, and wherein said ring optionally is substituted by F, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, oxo, CO(R₄₆), CON(R₄₇)(R₄₈), OH, N(R₅₀)CO(C₁-C₆)-alkyl or N(R₅₁)(R₅₂);

25 R₄₅ is OH, a 5- to 10-membered mono- or bicyclic ring which optionally contains one or two heteroatoms selected from the group consisting of N, O and S, wherein the 5- to 10-membered ring optionally is substituted by F, Cl, Br, OH, CF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₀-C₂)-alkylene-aryl, O-(C₀-C₂)-alkylene-aryl or N(R₅₁)(R₅₂).

30 4. The compound according to claim 3 wherein W is -CH=CH-.

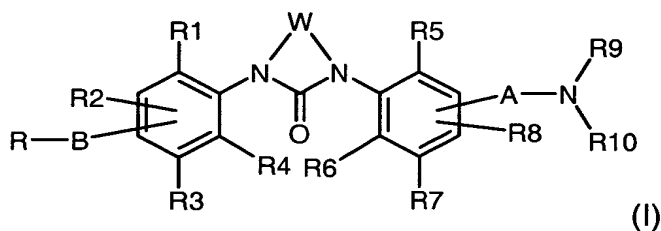
5. The compound according to claim 3 wherein m is 3 and R42, R43 and R44 are H.

6. The compound according to claim 3 wherein R5, R6, R7 and R8 are H.

7. The compound according to claim 3 wherein A and B are each disposed in the para position relative to the central W-containing heterocycle.

8. The compound according to claim 7 which is 1-[4-(2-dimethylaminoethoxy)-phenyl]-3-(4-phenoxyphenyl)-1,3-dihydroimidazol-2-one.

9. A method for the treatment or prevention of excessive weight or obesity in mammals comprising administering to said mammal a therapeutically effective amount of a compound, or a pharmaceutically acceptable salt thereof, of formula (I)



wherein

R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or spirocyclic ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R₁₁)(R₁₂), N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R₁₅)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ independently are H or (C₁-C₆)-alkyl;

B is a bond or a linker comprising one or two radicals selected from the group consisting of (C(R₁₉)(R₂₀))_i, C(OR₂₁)(R₂₂), O, N(R₂₃), S, SO, SO₂, and CO;

i is 1, 2 or 3;

R19, R20, R21, R22 and R23 independently are H, (C₁-C₆)-alkyl or aryl;

5 R1, R2, R3, and R4 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R24)(R25), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl,
10 CON(R26)(R27), N(R28)CO(R29), N(R30) SO₂(R31) or CO(R32);

R24, R25, R26, R27, R28 and R30 independently are H or (C₁-C₆)-alkyl;

R29, R31, and R32 independently are H, (C₁-C₆)-alkyl or aryl;

15

W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

n is 2, 3, 4 or 5;

20 R5, R6, R7 and R8 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R33)(R34), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl,
25 CON(R35)(R36), N(R37)CO(R38), N(R39) SO₂(R40), CO(R41) or a 5- to 7-membered heterocycle having 1 to 4 heteroatoms selected from the group consisting of O, N and S;

R33 and R34 independently are H or (C₁-C₆)-alkyl, or

30 R33 and R34 form together with the nitrogen atom to which they are bonded a 5- or 6-membered ring wherein when R33 and R34 form together with the nitrogen to

which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring optionally is O or S;

R35, R36, R37 and R39 independently are H or (C₁-C₆)-alkyl;

5

R38, R40 and R41 independently are H, (C₁-C₆)-alkyl or aryl;

A is a chain $-(C(R42)(R43))_m-$ wherein 0 to 2 members of the chain are optionally replaced by an element selected from the group consisting of O, S, N(R44), CO and SO₂;

10

m is 0, 1, 2, 3, 4 or 5;

R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

15

R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o-R45$, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl, CO-(CH₂)_oR45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl, CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 4 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

25

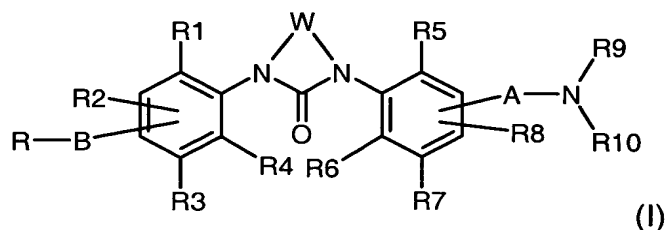
R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

30

o is 0, 1, 2, 3, 4, 5 or 6;

R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or more heteroatoms selected from the group consisting of N, O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-
 5 (C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R51)(R52), SO₂-CH₃ or COOH.

10. A method of treating a disease or psychiatric indication in mammals which
 10 comprises administering to said mammal a therapeutically effective amount of a compound, or a pharmaceutically acceptable salt thereof, of formula (I)



wherein

15 R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or spirocyclic ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally
 20 substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R11)(R12), N(R13)(R14), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R15)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

R11, R12, R13, R14 and R15 independently are H or (C₁-C₆)-alkyl;

25 B is a bond or a linker comprising one or two radicals selected from the group consisting of (C(R19)(R20))_i, C(OR21)(R22), O, N(R23), S, SO, SO₂, and CO;

i is 1, 2 or 3;

R19, R20, R21, R22 and R23 independently are H, (C₁-C₆)-alkyl or aryl;

R1, R2, R3, and R4 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R24)(R25), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R26)(R27), N(R28)CO(R29), N(R30) SO₂(R31) or CO(R32);

R24, R25, R26, R27, R28 and R30 independently are H or (C₁-C₆)-alkyl;

R29, R31, and R32 independently are H, (C₁-C₆)-alkyl or aryl;

W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

n is 2, 3, 4 or 5;

R5, R6, R7 and R8 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R33)(R34), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R35)(R36), N(R37)CO(R38), N(R39) SO₂(R40), CO(R41) or a 5- to 7-membered heterocycle having 1 to 4 heteroatoms selected from the group consisting of O, N and S;

R33 and R34 independently are H or (C₁-C₆)-alkyl, or

R33 and R34 form together with the nitrogen atom to which they are bonded a 5- or 6-membered ring wherein when R33 and R34 form together with the nitrogen to which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring optionally is O or S;

R35, R36, R37 and R39 independently are H or (C₁-C₆)-alkyl;

R38, R40 and R41 independently are H, (C₁-C₆)-alkyl or aryl;

5

A is a chain $-(C(R42)(R43))_m-$ wherein 0 to 2 members of the chain are optionally replaced by an element selected from the group consisting of O, S, N(R44), CO and SO₂;

10

m is 0, 1, 2, 3, 4 or 5;

R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

15

R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o-R45$, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl, CO-(CH₂)_oR45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl, CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

20

R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 4 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

25

R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

o is 0, 1, 2, 3, 4, 5 or 6;

30

R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or more heteroatoms selected from the group consisting of N,

O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R51)(R52), SO₂-CH₃ or COOH.

11. The method of claim 10 wherein the disease or psychiatric indication is selected from the group consisting of Type II diabetes, arteriosclerosis, high blood pressure, depression, anxiety, anxiety neuroses and schizophrenia.

12. The method of claim 11 wherein the disease is Type II diabetes.

13. The method of claim 11 wherein the disease is arteriosclerosis.

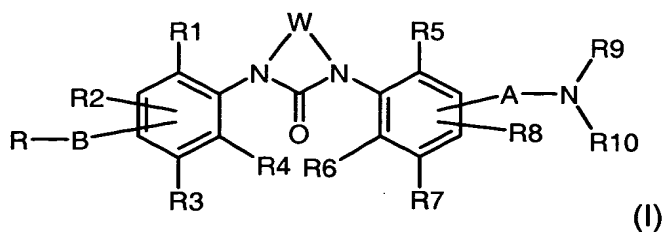
14. The method of claim 11 wherein the disease is high blood pressure.

15. The method of claim 11 wherein the psychiatric indication is depression.

16. The method of claim 11 wherein the psychiatric indication is anxiety or anxiety neuroses.

17. The method of claim 11 wherein the psychiatric indication is schizophrenia.

18. A method for the treatment or prevention of excessive weight or obesity in mammals comprising administering to said mammal a therapeutically effective amount of a mixture of one or more antiobesity agents or appetite-regulating active ingredients and a compound, or a pharmaceutically acceptable salt thereof, of formula (I)



wherein

R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-
 (C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or
 5 spirocyclic ring optionally containing one or more heteroatoms selected from the
 group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally
 substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R₁₁)(R₁₂),
 N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R₁₅)CO(C₁-C₆)-alkyl or
 COO-(C₁-C₆)-alkyl;

10 R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ independently are H or (C₁-C₆)-alkyl;

B is a bond or a linker comprising one or two radicals selected from the group
 consisting of (C(R₁₉)(R₂₀))_i, C(OR₂₁)(R₂₂), O, N(R₂₃), S, SO, SO₂, and CO;

15 i is 1, 2 or 3;

R₁₉, R₂₀, R₂₁, R₂₂ and R₂₃ independently are H, (C₁-C₆)-alkyl or aryl;

20 R₁, R₂, R₃, and R₄ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN,
 OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-
 alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-
 cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-
 C₈)-alkylene-aryl, S-aryl, N(R₂₄)(R₂₅), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl,
 25 CON(R₂₆)(R₂₇), N(R₂₈)CO(R₂₉), N(R₃₀) SO₂(R₃₁) or CO(R₃₂);

R₂₄, R₂₅, R₂₆, R₂₇, R₂₈ and R₃₀ independently are H or (C₁-C₆)-alkyl;

R₂₉, R₃₁, and R₃₂ independently are H, (C₁-C₆)-alkyl or aryl;

30 W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

n is 2, 3, 4 or 5;

R5, R6, R7 and R8 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN,
5 OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-
alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-
cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-
C₈)-alkylene-aryl, S-aryl, N(R33)(R34), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl,
10 CON(R35)(R36), N(R37)CO(R38), N(R39) SO₂(R40), CO(R41) or a 5- to 7-
membered heterocycle having 1 to 4 heteroatoms selected from the group consisting
of O, N and S;

R33 and R34 independently are H or (C₁-C₆)-alkyl, or
R33 and R34 form together with the nitrogen atom to which they are bonded a 5- or
15 6-membered ring wherein when R33 and R34 form together with the nitrogen to
which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring
optionally is O or S;

R35, R36, R37 and R39 independently are H or (C₁-C₆)-alkyl;

20

R38, R40 and R41 independently are H, (C₁-C₆)-alkyl or aryl;

A is a chain -(C(R42)(R43))_m- wherein 0 to 2 members of the chain are
optionally replaced by an element selected from the group consisting of O, S, N(R44),
25 CO and SO₂;

m is 0, 1, 2, 3, 4 or 5;

R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

30

R9 and R10 independently are H, (C₁-C₈)-alkyl, -(CH₂)_o-R45, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl, CO-(CH₂)_o-R45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl, CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

- 5 R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 4 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-
10 C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

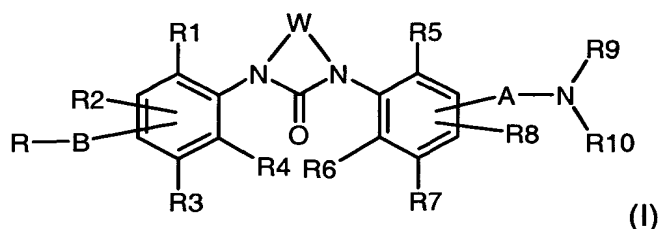
R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

- 15 o is 0, 1, 2, 3, 4, 5 or 6;

- R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or more heteroatoms selected from the group consisting of N, O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-
20 (C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R51)(R52), SO₂-CH₃ or COOH.

- 25 19. The method of claim 18 wherein the antiobesity agent or appetite-regulating active ingredient is selected from the group consisting of leptin, modified leptin, dexamphetamine, amphetamine, fenfluramine, dexfenfluramine, sibutramine, the mono- and bis-demethylated active metabolites of sibutramine, orlistat, mazindol, diethylpropion and phenteramine.

20. A method for the treatment or prevention of Type II diabetes in mammals comprising administering to said mammal a therapeutically effective amount of a mixture of one or more antidiabetics or hypoglycemic active ingredients and a compound, or a pharmaceutically acceptable salt thereof, of formula (I)



wherein

R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or spirocyclic ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R₁₁)(R₁₂), N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R₁₅)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ independently are H or (C₁-C₆)-alkyl;

B is a bond or a linker comprising one or two radicals selected from the group consisting of (C(R₁₉)(R₂₀))_i, C(OR₂₁)(R₂₂), O, N(R₂₃), S, SO, SO₂, and CO;

i is 1, 2 or 3;

R₁₉, R₂₀, R₂₁, R₂₂ and R₂₃ independently are H, (C₁-C₆)-alkyl or aryl;

R₁, R₂, R₃, and R₄ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-

C₈)-alkylene-aryl, S-aryl, N(R₂₄)(R₂₅), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₂₆)(R₂₇), N(R₂₈)CO(R₂₉), N(R₃₀) SO₂(R₃₁) or CO(R₃₂);

R₂₄, R₂₅, R₂₆, R₂₇, R₂₈ and R₃₀ independently are H or (C₁-C₆)-alkyl;

5

R₂₉, R₃₁, and R₃₂ independently are H, (C₁-C₆)-alkyl or aryl;

W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

10

n is 2, 3, 4 or 5;

15

R₅, R₆, R₇ and R₈ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R₃₃)(R₃₄), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₃₅)(R₃₆), N(R₃₇)CO(R₃₈), N(R₃₉) SO₂(R₄₀), CO(R₄₁) or a 5- to 7-membered heterocycle having 1 to 4 heteroatoms selected from the group consisting of O, N and S;

20

R₃₃ and R₃₄ independently are H or (C₁-C₆)-alkyl, or

R₃₃ and R₃₄ form together with the nitrogen atom to which they are bonded a 5- or 6-membered ring wherein when R₃₃ and R₃₄ form together with the nitrogen to which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring optionally is O or S;

25

R₃₅, R₃₆, R₃₇ and R₃₉ independently are H or (C₁-C₆)-alkyl;

R₃₈, R₄₀ and R₄₁ independently are H, (C₁-C₆)-alkyl or aryl;

30

A is a chain $-(C(R42)(R43))_m-$ wherein 0 to 2 members of the chain are optionally replaced by an element selected from the group consisting of O, S, N(R44), CO and SO₂;

5 m is 0, 1, 2, 3, 4 or 5;

R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

10 R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o-R45$, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-alkyl, CO-(CH₂)_oR45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl, CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

15 R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to 10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen atom, may optionally contain 0 to 4 additional heteroatoms selected from the group consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br, CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

20

R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

o is 0, 1, 2, 3, 4, 5 or 6;

25

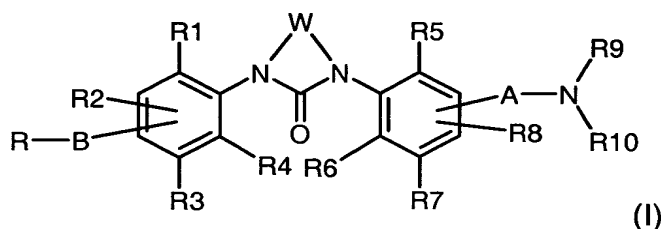
R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which optionally contains one or more heteroatoms selected from the group consisting of N, O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R51)(R52), SO₂-CH₃ or COOH.

30

21. The method of claim 20 wherein said antidiabetic or hypoglycemic active ingredient is selected from the group consisting of insulin, a sulfonylurea, a biguanide, a meglitinide, a thiazolidinedione, an oxadiazolidinedione and an α -glucosidase inhibitor.

5

22. A pharmaceutical composition comprising a pharmaceutical carrier and a therapeutically effective amount of a compound, or a pharmaceutically acceptable salt thereof, of formula (I)



10 wherein

R is (C₁-C₈)-alkyl, (C₀-C₈)-alkylene-aryl, (C₃-C₈)-cycloalkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₂-C₈)-alkenyl, (C₂-C₈)-alkynyl, a 3- to 12-membered mono-, bi- or spirocyclic ring optionally containing one or more heteroatoms selected from the group consisting of N, O, and S, and wherein the 3- to 12-membered ring is optionally substituted by F, Cl, Br, NO₂, CF₃, OCF₃, CN, (C₁-C₆)-alkyl, aryl, CON(R₁₁)(R₁₂), N(R₁₃)(R₁₄), OH, O-(C₁-C₆)-alkyl, S-(C₁-C₆)-alkyl, N(R₁₅)CO(C₁-C₆)-alkyl or COO-(C₁-C₆)-alkyl;

20 R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ independently are H or (C₁-C₆)-alkyl;

B is a bond or a linker comprising one or two radicals selected from the group consisting of (C(R₁₉)(R₂₀))_i, C(OR₂₁)(R₂₂), O, N(R₂₃), S, SO, SO₂, and CO;

25 i is 1, 2 or 3;

R₁₉, R₂₀, R₂₁, R₂₂ and R₂₃ independently are H, (C₁-C₆)-alkyl or aryl;

R1, R2, R3, and R4 independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R₂₄)(R₂₅), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₂₆)(R₂₇), N(R₂₈)CO(R₂₉), N(R₃₀) SO₂(R₃₁) or CO(R₃₂);

R₂₄, R₂₅, R₂₆, R₂₇, R₂₈ and R₃₀ independently are H or (C₁-C₆)-alkyl;

R₂₉, R₃₁, and R₃₂ independently are H, (C₁-C₆)-alkyl or aryl;

W is -(CH₂)_n-, -CH=CH-, -CH=N- or -N=CH-;

n is 2, 3, 4 or 5;

R₅, R₆, R₇ and R₈ independently are H, F, Cl, Br, I, OH, CF₃, NO₂, CN, OCF₃, O-(C₁-C₆)-alkyl, O-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R₃₃)(R₃₄), SO₂-CH₃, COOH, COO-(C₁-C₆)-alkyl, CON(R₃₅)(R₃₆), N(R₃₇)CO(R₃₈), N(R₃₉) SO₂(R₄₀), CO(R₄₁) or a 5- to 7-membered heterocycle having 1 to 4 heteroatoms selected from the group consisting of O, N and S;

R₃₃ and R₃₄ independently are H or (C₁-C₆)-alkyl, or

R₃₃ and R₃₄ form together with the nitrogen atom to which they are bonded a 5- or 6-membered ring wherein when R₃₃ and R₃₄ form together with the nitrogen to which they are bonded a 6-membered ring, one CH₂ group of the 6-membered ring optionally is O or S;

R₃₅, R₃₆, R₃₇ and R₃₉ independently are H or (C₁-C₆)-alkyl;

R38, R40 and R41 independently are H, (C₁-C₆)-alkyl or aryl;

A is a chain $-(C(R42)(R43))_m-$ wherein 0 to 2 members of the chain are
5 optionally replaced by an element selected from the group consisting of O, S, N(R44), CO and SO₂;

m is 0, 1, 2, 3, 4 or 5;

10 R42, R43, R44 independently are H, (C₁-C₆)-alkyl or aryl;

R9 and R10 independently are H, (C₁-C₈)-alkyl, $-(CH_2)_o-R45$, (C₁-C₄)-alkoxy-
(C₁-C₄)-alkyl, aryloxy-(C₁-C₄)-alkyl, (C₃-C₈)-alkenyl, (C₃-C₈)-alkynyl, CO-(C₁-C₈)-
alkyl, CO-(CH₂)_oR45, CO-(C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, CO-aryloxy-(C₁-C₄)-alkyl,
15 CO-(C₂-C₈)-alkenyl, CO-(C₂-C₈)-alkynyl, or

R9 and R10 form together with the nitrogen atom to which they are bonded a 4- to
10-membered mono-, bi- or spirocyclic ring wherein said ring, apart from the nitrogen
atom, may optionally contain 0 to 4 additional heteroatoms selected from the group
consisting of O, N and S and wherein said ring optionally is substituted by F, Cl, Br,
20 CF₃, NO₂, CN, (C₁-C₆)-alkyl, O-(C₁-C₈)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, (C₀-
C₈)-alkylene-aryl, oxo, CO(R46), CON(R47)(R48), OH, COO(R49), N(R50)CO(C₁-
C₆)-alkyl, N(R51)(R52) or SO₂CH₃;

R46, R47, R48, R49, R50, R51 and R52 independently are H or (C₁-C₄)-alkyl;

25 o is 0, 1, 2, 3, 4, 5 or 6;

R45 is OH, CH(aryl)₂, a 3- to 12-membered mono- or bicyclic ring which
optionally contains one or more heteroatoms selected from the group consisting of N,
30 O and S wherein the 3- to 12-membered ring optionally is substituted by F, Cl, Br, I,
OH, CF₃, NO₂, CN, OCF₃, oxo, O-(C₁-C₆)-alkyl, (C₁-C₄)-alkoxy-(C₁-C₄)-alkyl, S-

(C₁-C₆)-alkyl, (C₁-C₆)-alkyl, (C₂-C₆)-alkenyl, (C₃-C₈)-cycloalkyl, O-(C₃-C₈)-cycloalkyl, (C₃-C₈)-cycloalkenyl, O-(C₃-C₈)-cycloalkenyl, (C₂-C₆)-alkynyl, (C₀-C₈)-alkylene-aryl, O-(C₀-C₈)-alkylene-aryl, S-aryl, N(R₅₁)(R₅₂), SO₂-CH₃ or COOH.

- 5 23. A pharmaceutical composition according to claim 22 further comprising a therapeutically effective amount of one or more compounds selected from the group consisting of an antiobesity agent, an appetite-regulating active ingredient, an antidiabetic and a hypoglycemic active ingredient.
- 10 24. A pharmaceutical composition according to claim 23 wherein said antiobesity agent or appetite-regulating active ingredient is selected from the group consisting of of leptin, modified leptin, dexamphetamine, amphetamine, fenfluramine, dexfenfluramine, sibutramine, the mono- and bis-demethylated active metabolites of sibutramine, orlistat, mazindol, diethylpropion and phenteramine.
- 15 25. A pharmaceutical composition according to claim 23 wherein said antidiabetic or hypoglycemic active ingredient is selected from the group consisting of insulin, a sulfonylurea, a biguanide, a meglitinide, a thiazolidinedione, an oxadiazolidinedione and an α -glucosidase inhibitor.